## **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions and listings of claims in the application:

## LISTING OF CLAIMS:

- 1. (currently amended): An Her-2/neu plasmid construct having anti-cancer activity which is prepared by inserting A pTV2 or pCK vector comprising a nucleotide sequence encoding a truncated human Her-2/neu geneprotein, said truncated human Her-2/neu protein lacking the an intracellular domain-into plasmid pTV2 or pCK.
- 2. (currently amended): The <u>plasmid construct\_vector</u> of claim 1, wherein the human Her-2/neu gene has the nucleotide sequence of SEQ ID NO: 2.
- 3. (currently amended): The plasmid construct vector of claim 2, which is pNeu<sub>TM</sub> (KCCM-10393) or pCK<sub>TM</sub> (KCCM-10396).
- 4. (withdrawn) The plasmid construct of claim 1, whrein the truncated human Her-2/neu gene further lacks the transmembrane domain.
- 5. (withdrawn) The plasmid construct of claim 4, wherein the human Her-2/neu gene has the nucleotide of SEQ ID NO: 3.
- 6. (withdrawn) The plasmid construct of claim 5, which is pNeu<sub>ECD</sub> (KCCM-10394) or pCK<sub>ECD</sub> (KCCM-10395).
- 7. (withdrawn) The plasmid construct of claim 1, wherein the signal peptide of the human Her-2/neu gene is replaced by the signal peptide of herpes simplex type I glycoprotein D (gD).

- 8. (withdrawn) The plasmid construct of claim 7, which is pNeu<sub>TM-gDs</sub>.
- 9. (withdrawn) The plasmid construct of claim 4, wherein the signal peptide of the human Her-2/neu gene is replaced by the signal peptide of herpes simplex type I glycoprotein D (gD).
  - 10. (withdrawn) The plasmid construct of claim 7, which is pNeu<sub>ECD-gDs</sub>.
- 11. (currently amended): The <u>plasmid construct\_vector</u> of claim 1, which further translates a cytokine gene besides the human Her 2/neu gene comprises a nucleotide sequence encoding a cytokine.
- 12. (currently amended): The <u>plasmid construct vector</u> of claim 11, wherein the cytokine gene is <u>selected from the group consisting of granulocyte-macrophage colony-stimulating factor (GM-CSF), FMS-like tyrosine kinase 3 ligand (Flt3L), early T-lymphocyte activation 1 (Eta-1), interleukin 12 (IL-12), IL-15 and IL-18.</u>
- 13. (currently amended): A DNA vaccine <u>composition</u> for preventing and/or treating cancer, which comprises the plasmid construct of claim 1 as an effective ingredient and a pharmaceutically acceptable carrier. comprising a pTV2 vector or pCK vector which comprises a nucleotide sequence encoding a truncated human Her-2/neu protein, said truncated human Her-2/neu protein lacking an intracellular domain.
- 14. (currently amended): The DNA vaccine <u>composition</u> of claim 13, which further comprises a <u>cytokine gene expressing</u>-plasmid <u>which expresses a gene encoding a cytokine</u>.
- 15. (currently amended): The DNA vaccine <u>composition</u> of claim 14, wherein the cytokine <del>gene</del> is <del>selected from the group consisting of GM-CSF, Flt3L, Eta-1, IL-12, IL-15 and IL-18</del>.

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- 16. (currently amended): A method for preventing and/or treating cancer, which comprises the step of administering an effective amount of the DNA vaccine <u>composition</u> of claim 13.
- 17. (new): The DNA vaccine composition of claim 13, wherein the pTV2 vector or pCK vector further comprises a nucleotide sequence encoding a cytokine.
- 18. (new): The DNA vaccine composition of claim 17, wherein the cytokine is GM-CSF.